# familial male-limited precocious puberty

Familial male-limited precocious puberty is a condition that causes early sexual development in males; females are not affected. Boys with this disorder begin exhibiting the signs of puberty in early childhood, between the ages of 2 and 5. Signs of male puberty include a deepening voice, acne, increased body hair, underarm odor, growth of the penis and testes, and spontaneous erections. Changes in behavior, such as increased aggression and early interest in sex, may also occur. Without treatment, affected boys grow quickly at first, but they stop growing earlier than usual. As a result, they tend to be shorter in adulthood compared with other members of their family.

# Frequency

Familial male-limited precocious puberty is a rare disorder; its prevalence is unknown.

## **Genetic Changes**

Familial male-limited precocious puberty can be caused by mutations in the *LHCGR* gene. This gene provides instructions for making a receptor protein called the luteinizing hormone/chorionic gonadotropin receptor. Receptor proteins have specific sites into which certain other proteins, called ligands, fit like keys into locks. Together, ligands and their receptors trigger signals that affect cell development and function.

The protein produced from the *LHCGR* gene acts as a receptor for two ligands: luteinizing hormone and a similar hormone called chorionic gonadotropin. The receptor allows the body to respond appropriately to these hormones. In males, chorionic gonadotropin stimulates the development of cells in the testes called Leydig cells, and luteinizing hormone triggers these cells to produce androgens. Androgens, including testosterone, are the hormones that control male sexual development and reproduction. In females, luteinizing hormone triggers the release of egg cells from the ovaries (ovulation); chorionic gonadotropin is produced during pregnancy and helps maintain conditions necessary for the pregnancy to continue.

Certain *LHCGR* gene mutations result in a receptor protein that is constantly turned on (constitutively activated), even when not attached (bound) to luteinizing hormone or chorionic gonadotropin. In males, the overactive receptor causes excess production of testosterone, which triggers male sexual development and lead to early puberty in affected individuals. The overactive receptor has no apparent effect on females.

Approximately 18 percent of individuals with familial male-limited precocious puberty have no identified *LHCGR* gene mutation. In these individuals, the cause of the disorder is unknown.

#### Inheritance Pattern

This condition is inherited in an autosomal dominant, male-limited pattern, which means one copy of the altered *LHCGR* gene in each cell is sufficient to cause the disorder in males. Females with mutations associated with familial male-limited precocious puberty appear to be unaffected. In some cases, an affected male inherits the mutation from either his mother or his father. Other cases result from new mutations in the gene and occur in males with no history of the disorder in their family.

#### Other Names for This Condition

- familial gonadotrophin-independent sexual precocity
- GIPP
- gonadotrophin-independent precocious puberty
- precocious pseudopuberty
- pubertas praecox
- testotoxicosis

## **Diagnosis & Management**

## **Genetic Testing**

 Genetic Testing Registry: Gonadotropin-independent familial sexual precocity https://www.ncbi.nlm.nih.gov/gtr/conditions/C0342549/

# Other Diagnosis and Management Resources

 Boston Children's Hospital: Precocious Puberty http://www.childrenshospital.org/conditions-and-treatments/conditions/precocious-early-puberty

#### General Information from MedlinePlus

- Diagnostic Tests
   https://medlineplus.gov/diagnostictests.html
- Drug Therapy https://medlineplus.gov/drugtherapy.html
- Genetic Counseling https://medlineplus.gov/geneticcounseling.html
- Palliative Care https://medlineplus.gov/palliativecare.html
- Surgery and Rehabilitation https://medlineplus.gov/surgeryandrehabilitation.html

#### **Additional Information & Resources**

#### MedlinePlus

- Encyclopedia: Precocious Puberty https://medlineplus.gov/ency/article/001168.htm
- Health Topic: Endocrine Diseases https://medlineplus.gov/endocrinediseases.html
- Health Topic: Puberty https://medlineplus.gov/puberty.html

## Genetic and Rare Diseases Information Center

- Precocious puberty https://rarediseases.info.nih.gov/diseases/7446/precocious-puberty
- Testotoxicosis https://rarediseases.info.nih.gov/diseases/4475/testotoxicosis

#### Additional NIH Resources

 Eunice Kennedy Shriver National Institute of Child Health and Human Development: Precocious Puberty https://www.nichd.nih.gov/health/topics/puberty/Pages/default.aspx

#### **Educational Resources**

- Disease InfoSearch: Testotoxicosis http://www.diseaseinfosearch.org/Testotoxicosis/7032
- KidsHealth: Precocious Puberty http://kidshealth.org/en/parents/precocious.html
- MalaCards: familial male-limited precocious puberty http://www.malacards.org/card/familial\_male\_limited\_precocious\_puberty
- Orphanet: Familial male-limited precocious puberty http://www.orpha.net/consor/cgi-bin/OC\_Exp.php?Lng=EN&Expert=3000
- University of Michigan Health System: Precocious Puberty http://www.med.umich.edu/yourchild/topics/puberty.htm

## Patient Support and Advocacy Resources

- Magic Foundation for Children's Growth https://www.magicfoundation.org/Growth-Disorders/Precocious-Puberty/
- National Organization for Rare Disorders: Precocious Puberty https://rarediseases.org/rare-diseases/precocious-puberty/

# ClinicalTrials.gov

ClinicalTrials.gov
 https://clinicaltrials.gov/ct2/results?cond=%22familial+male-limited+precocious
 +puberty%22+OR+%22Puberty%2C+Precocious%22

#### Scientific Articles on PubMed

PubMed

https://www.ncbi.nlm.nih.gov/pubmed?term=%28Puberty,+Precocious%5BMAJ R%5D%29+AND+%28%28familial+male-limited+precocious+puberty%5BTIAB %5D%29+OR+%28testotoxicosis%5BTIAB%5D%29%29+AND+english%5Bla%5D +AND+human%5Bmh%5D+AND+%22last+3600+days%22%5Bdp%5D

#### OMIM

 PRECOCIOUS PUBERTY, MALE-LIMITED http://omim.org/entry/176410

## **Sources for This Summary**

- Brito VN, Latronico AC, Arnhold IJ, Mendonça BB. Update on the etiology, diagnosis and therapeutic management of sexual precocity. Arq Bras Endocrinol Metabol. 2008 Feb;52(1):18-31. Review. Erratum in: Arq Bras Endocrinol Metabol. 2008 Apr;52(3):576. Citation on PubMed: https://www.ncbi.nlm.nih.gov/pubmed/18345393
- Chan WY. Disorders of sexual development caused by luteinizing hormone receptor mutations. Beijing Da Xue Xue Bao. 2005 Feb 18;37(1):32-8.
   Citation on PubMed: https://www.ncbi.nlm.nih.gov/pubmed/15719037
- Soriano-Guillen L, Mitchell V, Carel JC, Barbet P, Roger M, Lahlou N. Activating mutations in the luteinizing hormone receptor gene: a human model of non-follicle-stimulating hormone-dependent inhibin production and germ cell maturation. J Clin Endocrinol Metab. 2006 Aug;91(8):3041-7. Epub 2006 May 9.
  - Citation on PubMed: https://www.ncbi.nlm.nih.gov/pubmed/16684832
- Themmen AP, Verhoef-Post M. LH receptor defects. Semin Reprod Med. 2002 Aug;20(3):199-204.
   Review.
  - Citation on PubMed: https://www.ncbi.nlm.nih.gov/pubmed/12428200
- Traggiai C, Stanhope R. Disorders of pubertal development. Best Pract Res Clin Obstet Gynaecol. 2003 Feb;17(1):41-56. Review.
   Citation on PubMed: https://www.ncbi.nlm.nih.gov/pubmed/12758225

## Reprinted from Genetics Home Reference:

https://ghr.nlm.nih.gov/condition/familial-male-limited-precocious-puberty

Reviewed: August 2012 Published: March 21, 2017 Lister Hill National Center for Biomedical Communications U.S. National Library of Medicine National Institutes of Health Department of Health & Human Services